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DEBRIS CHARGE STATES IN HANE (HIGH ALTITUDE NUCLEAR  
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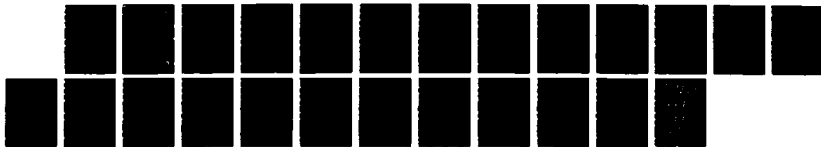
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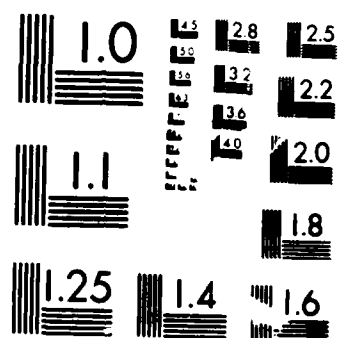
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NRL Memorandum Report 6085

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## Debris Charge States in HANE and in the NRL Laser Experiment

E. HYMAN

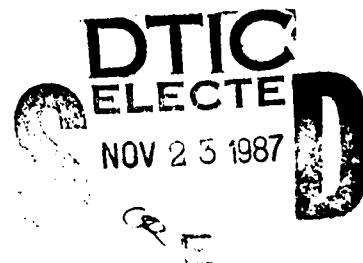
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October 30, 1987

This report was sponsored by DNA under "Weapons Phenomenology and Code Development"  
Work Unit Code & Title: RB RC/00158, Plasma Structure Evolution.



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AD-A187541

SECURITY CLASSIFICATION OF THIS PAGE

## REPORT DOCUMENTATION PAGE

|   |   |   |  |
|---|---|---|--|
| 1a. REPORT SECURITY CLASSIFICATION<br>UNCLASSIFIED  |   | 1b. RESTRICTIVE MARKINGS  |  |
| 2a. SECURITY CLASSIFICATION AUTHORITY   |   | 3. DISTRIBUTION/AVAILABILITY OF REPORT<br>Approved for public release;<br>distribution unlimited. |  |
| 2b. DECLASSIFICATION/DOWNGRADING SCHEDULE   |   |   |  |
| 4. PERFORMING ORGANIZATION REPORT NUMBER(S)<br>NRL Memorandum Report 6085   |   | 5. MONITORING ORGANIZATION REPORT NUMBER(S)   |  |
| 6a. NAME OF PERFORMING ORGANIZATION<br>Naval Research Laboratory  | 6b. OFFICE SYMBOL<br>(If applicable)<br>Code 4780 | 7a. NAME OF MONITORING ORGANIZATION   |  |
| 6c. ADDRESS (City, State, and ZIP Code)<br>Washington, DC 20375-5000  |   | 7b. ADDRESS (City, State, and ZIP Code)   |  |
| 8a. NAME OF FUNDING/SPONSORING ORGANIZATION<br>Defense Nuclear Agency   | 8b. OFFICE SYMBOL<br>(If applicable)<br>RAAE      | 9. PROCUREMENT INSTRUMENT IDENTIFICATION NUMBER<br>47-0889-07 (MIPR No. 87-518)                   |  |
| 8c. ADDRESS (City, State, and ZIP Code)<br>Washington, DC 20305   |   | 10. SOURCE OF FUNDING NUMBERS   |  |
|   |   | PROGRAM<br>ELEMENT NO.<br>62715H  | PROJECT<br>NO.<br>S99QMXBC -<br>00102<br>WORK UNIT<br>ACCESSION NO.<br>DN580-072 |
| 11. TITLE (Include Security Classification)<br>Debris Charge States in Hane and in the NRL Laser Experiment   |   |   |  |
| 12. PERSONAL AUTHOR(S)<br>Hyman*, E., Mulbrandon, M. and Giuliani, J.L., Jr.  |   |   |  |
| 13a. TYPE OF REPORT<br>Interim  | 13b. TIME COVERED<br>FROM TO                      | 14. DATE OF REPORT (Year, Month, Day)<br>1987 October 30  | 15. PAGE COUNT<br>23   |
| 16. SUPPLEMENTARY NOTATION<br>*This report was sponsored by DNA under "Weapons Phenomenology and Code Development."<br>Work Unit Code and Title: RB RC/00158, Plasma Structure Evolution.   |   |   |  |
| 17. COSATI CODES  |   | 18. SUBJECT TERMS (Continue on reverse if necessary and identify by block number)                 |  |
| FIELD   | GROUP   | SUB-GROUP   |  |
|   |   | HANE (High Altitude Nuclear Explosion)  |  |
|   |   | NRL Laser Experiment, Debris Charge State,  |  |
|   |   | Scaling, Recombination Rates  |  |
| 19. ABSTRACT (Continue on reverse if necessary and identify by block number)  |   |   |  |
| <p>We show that in the NRL laser experiment the expanding debris ions remain highly charged, in contrast to the situation in a high altitude nuclear explosion (HANE) in which recombination to a low charge state is rapid. The difference in the two cases is a result of (1) a failure in the scaling of debris density relative to a HANE in the early expansion phase and (2) a failure in scaling of the energy deposition time. Results based on simple idealized models for the HANE and laser experiment are presented and these are compared with more sophisticated calculations.</p> <p style="text-align: right;">(Keegan)</p> |   |   |  |
| 20. DISTRIBUTION/AVAILABILITY OF ABSTRACT<br><input checked="" type="checkbox"/> UNCLASSIFIED/UNLIMITED <input type="checkbox"/> SAME AS RPT. <input type="checkbox"/> DTIC USERS   |   | 21. ABSTRACT SECURITY CLASSIFICATION<br>UNCLASSIFIED  |  |
| 22a. NAME OF RESPONSIBLE INDIVIDUAL<br>J.D. HUBA  |   | 22b. TELEPHONE (Include Area Code)<br>202-767-3630  | 22c. OFFICE SYMBOL<br>Code 4780  |

DD FORM 1473, 84 MAR

83 APR edition may be used until exhausted  
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SECURITY CLASSIFICATION OF THIS PAGE

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## DEBRIS CHARGE STATES IN HANE AND IN THE NRL LASER EXPERIMENT

### I. Introduction

It is reasonably well established that the charge state,  $z$ , of HANE debris drops from very high values to  $z$  close to 1 on disassembly time scales. Both observations and disassembly calculations suggest this result. A calculation by Clark and Jacobs at NRL<sup>(1)</sup> estimates that the charge state drops to  $z \sim 1$  by the time the burst has expanded to  $\sim 200$  meters radius, at which point the charge state is frozen in. That is, by this time the density has dropped to the point that recombination is too slow to be important and, further, future temperature decreases will not be sufficient to increase the recombination coefficient enough to offset the density decrease.

At the Naval Research Laboratory a laser target experiment, PHAROS II<sup>(2)</sup>, has been designed and operated to model on a reduced scale some of the physics occurring in an actual HANE. Simulations of the laser experiment using a hydro-chemistry-radiation code HANEX<sup>(3)</sup> predict very high charge states ( $z \sim 10$ ) for the laser experiment in the forward moving debris, which persist, at least, until interaction with the background gas becomes important. We have identified two critical differences between the laser experiment and the HANE event that are responsible for the disparity in the results:

- (1) a failure in density scaling in the adiabatic expansion phase, and
- (2) a difference in scaled times for the deposition of the laser/nuclear energy, during disassembly.

The laser experiment was designed to scale relative to HANE in the following way,<sup>(4)</sup>

$$\begin{aligned} t_H &= 10^6 t_L \\ r_H &= 10^6 r_L \\ n_H &= 10^{-6} n_L \\ M_H &= 10^{12} M_L \\ E_H &= 10^{12} E_L \end{aligned} \tag{1}$$

where  $t$ ,  $r$ ,  $n$ ,  $M$ , and  $E$  are respectively time, expansion radius, background density, debris mass, and energy for HANE (H) and the laser experiment (L). It follows from Eq. 1, that 2-body reaction rate time scales should be related by  $\tau_H = 10^6 \tau_L$  (since  $\tau = (n\alpha)^{-1}$ , where  $\alpha$  is the rate coefficient).

In this note, we will show that there is a breakdown in scaling, and that recombination is much less effective in the laser experiment than it is in HANE, with the result that the laser debris remains highly charged. In Section II we will show that during the debris expansion, recombination times are the same in HANE and in the laser experiment (they do not scale, as suggested by Eq. 1). Also, "problem time" scales like  $10^4$ , not  $10^6$ , during the expansion, as we will see. In Section III we introduce an idealized HANE model and an idealized model for the laser experiment to clarify the essential distinctions. In Section IV we compare these results with HANEX code calculations and present our conclusions.

## II. Scaling and Debris Recombination

To evaluate the importance of recombination (or ionization) in reducing (increasing) the average charge state of the debris, it is convenient to introduce the dimensionless parameter,  $\xi$ , the fraction of ions that will recombine or ionize in a time interval  $\delta t$

$$\xi = \frac{\delta t}{\tau} \tag{2}$$

where  $\tau = (N_e \alpha)^{-1}$ , ( $N_e$  = electron density) is the recombination (ionization) time. In this discussion we neglect 3-body recombination, which is unimportant at the high temperatures associated with the early debris expansion. It is included in our model, however. We choose  $\delta t$  to be the time scale for changes in the density and temperature (an expansion time scale). Then the expanding plasma charge state will be in equilibrium if  $\xi$  for both recombination and ionization are large. If  $\xi$  becomes less than unity the plasma may no longer be in equilibrium. In particular, if  $\xi$  for recombination processes becomes and stays much less than unity, then that charge state will be frozen at its existing value, i.e., recombination will not occur.

If the debris expansion scaling satisfied Eq. 1, then according to Eq. 2,  $\xi$  for HANE and the laser experiment would be the same. However, this is not the case for debris recombination during the early expansion of the debris. Here, the ambient density is negligible compared to the debris density. Both in the laser experiment and in the HANE the debris density starts from the solid state. Thus, debris density is not scaled at all in the expansion phase; it is the same for HANE and the laser.

If the laser experiment were strictly a scaled down HANE (mass scaled by  $10^{-12}$ ) with the same initial temperature, the time scale for recombination (and ionization),  $\tau$ , would be identical. However, if mass ( $M$ ) scales  $\sim 10^{12}$ , and because debris densities are the same, the expansion radius scales  $\sim (M)^{1/3} \sim 10^4$ . Since expansion velocities are comparable, time ( $\delta t$ ) scales  $\sim 10^4$ , also. Then, from Eq. 2, for HANE  $\xi \sim 10^4$  times larger than in a laser experiment size HANE.

In the real experiment the temperature of the debris never exceeds  $10^3$  eV, while in HANE the temperature is greater than  $10^4$  eV at early times. If the laser stayed cooler than in a HANE, increased recombination at lower temperatures might compensate somewhat for the  $\xi$  discrepancy. However, in



the HANE event, the energy is released and deposited in a time  $\sim 10^{-6}$  sec. In the laser, scaling of problem time by  $10^{-4}$  implies an appropriate deposition time  $\sim 10^{-10}$  sec, but the actual deposition time is  $\sim 10^{-8}$  sec. Thus, there is a scaling discrepancy of order  $10^{-2}$ . In other words, the laser energy is deposited over a time period a factor of  $10^2$  too long compared to a HANE. Thus, while the HANE debris is expanding adiabatically, and cooling accordingly, the laser debris expands but is maintained at a high temperature as long as the laser is on.<sup>(3)</sup> Thereafter, the debris expands and cools adiabatically but the temperature remains higher than in a HANE, at the same densities. This higher temperature also decreases recombination processes in the laser experiment.

Thus, recombination is less effective in the laser experiment, first, because of scaling which requires  $\xi_R(\text{HANE}) = 10^4 \xi_R(\text{LASER})$  and, second, because the long laser energy deposition time delays cooling. In the next section we illustrate the result of these effects using simplified models for the HANE and the laser experiment.

### III. Results

The recombination coefficient is made up of 3 parts: radiative recombination, dielectronic recombination, and 3-body recombination. Their general dependence with temperature is quite different and explains their relative importance in different regimes. At intermediate temperatures dielectronic recombination tends to be larger than radiative and 3-body is unimportant. Dielectronic falls off somewhat faster than radiative ( $\sim T_e^{-3/2}$  vs.  $\sim T_e^{-1}$ ) with increasing temperature. At lower temperatures dielectronic drops off exponentially while radiative continues to increase. At very low temperatures 3-body will dominate even at low densities because it increases like  $T_e^{-9/2}$  as  $T_e$  decreases.<sup>(5)</sup> In Appendix I we present expressions for each of these rate coefficients.

Table 1 shows "typical" HANE parameters for ideal adiabatic expansion of aluminum into a vacuum. The plasma expands radially at constant velocity  $v = 1 \times 10^8$  cm/sec, according to

$$n = n_0 (r_0/R)^3; \quad (3)$$

where

$$R = vt$$

Assuming adiabatic expansion with  $\gamma = 5/3$ ,

$$T = T_0 (r_0/R)^2 \quad (4)$$

$T_0$ ,  $M_0$  and  $r_0$  are the temperature, density, and radius when the expansion begins. Included in the Table are temperature (T), debris ion density (n),

TABLE 1

| T<br>(ev) | $n$<br>( $\text{cm}^{-3}$ ) | t<br>(sec) | R<br>(cm) | $\xi_{R3}$ | $\xi_{RR}$ | $\xi_{RD}$ | $\xi_I$ | z  |
|-----------|-----------------------------|------------|-----------|------------|------------|------------|---------|----|
| 1.0E+05   | 1.0E+23                     | 1.0E-06    | 1.0E+02   | 0.0E+00    | 5.5E+04    | -----      | 1.2E+07 | 13 |
| 2.2E+04   | 1.0E+22                     | 2.2E-06    | 2.2E+02   | 0.0E+00    | 3.8E+04    | -----      | 4.0E+06 | 13 |
| 4.6E+03   | 1.0E+21                     | 4.6E-06    | 4.6E+02   | 7.4E+00    | 2.6E+04    | -----      | 6.7E+05 | 13 |
| 1.0E+03   | 1.0E+20                     | 1.0E-05    | 1.0E+03   | 2.3E+00    | 1.1E+04    | 2.5E+04    | 4.7E+04 | 12 |
| 2.2E+02   | 1.0E+19                     | 2.2E-05    | 2.2E+03   | 1.7E+00    | 2.4E+03    | 4.5E+04    | 5.1E+04 | 10 |
| 4.6E+01   | 1.0E+18                     | 4.6E-05    | 4.6E+03   | 2.7E+00    | 4.5E+02    | 1.3E+03    | 5.9E+03 | 6  |
| 1.0E+01   | 1.0E+17                     | 1.0E-04    | 1.0E+04   | 1.9E+01    | 6.4E+01    | 5.0E+01    | 5.6E-01 | 4  |
| 2.2E+00   | 1.0E+16                     | 2.2E-04    | 2.2E+04   | 9.2E+00    | 5.3E+00    | 9.1E+02    | 6.5E+01 | 2  |
| 4.6E-01   | 1.0E+15                     | 4.6E-04    | 4.6E+04   | 8.5E+00    | 3.2E-01    | 2.0E+01    | 7.8E-01 | 1  |
| 1.0E-01   | 1.0E+14                     | 1.0E-03    | 1.0E+05   | 4.4E+02    | 2.3E-01    | 6.3E-06    | 5.9E-18 | 1  |
| 2.2E-02   | 1.0E+13                     | 2.2E-03    | 2.2E+05   | 4.9E+03    | 1.7E-01    | 0.0E+00    | 5.9E-19 | 1  |

time ( $t$ ), and radial expansion ( $R$ ). We have calculated the parameter,  $\xi$ , for 3-body recombination ( $\xi_{R3}$ ), radiative recombination ( $\xi_{RR}$ ), dielectronic recombination ( $\xi_{RD}$ ), and ionization ( $\xi_I$ ). The last column is the approximate value of the dominant charge state,  $z$ . In the calculation of  $\xi$  we have approximated the expansion time scale  $\delta t$ , by the problem time  $t$ , given in the Table. The parameters  $\xi$  are calculated assuming recombination from  $z$  to  $z - 1$  and ionization from  $z - 1$  to  $z$ . In a cooling plasma if at least one of the  $\xi$ 's for recombination is greater than unity the plasma is in equilibrium and the charge state,  $z$ , is determined by the temperature,  $T$ .

At very early times the plasma is in equilibrium and is stripped. (Dielectronic recombination is not defined for a stripped ion.) As the adiabatic expansion continues and density and temperature drop, the plasma remains in equilibrium and the charge state begins to drop. Both dielectronic recombination and radiative recombination are important. Eventually, the temperature is sufficiently low that the charge state drops to  $z \sim 1$  ( $n \sim 10^{15} \text{ cm}^{-3}$ ). At this time dielectronic and 3 body recombination are comparable. As density and temperature are reduced further ( $n < 10^{15} \text{ cm}^{-3}$ ) 3-body recombination begins to dominate. Here, we need to stress that we have used a model of an ideal adiabatic expansion. In the real case, each recombination releases energy, much of which ends up heating the electrons. Thus, the plasma will no longer cool adiabatically and the temperature will not drop so precipitously. As the density continues to decrease below  $\sim 10^{15} \text{ cm}^{-3}$ , the debris is likely to be frozen at  $z$  close to 1.

The effect of the failure of density scaling during the expansion can be made clear by scaling the HANE expansion of Table 1 to the size of a laser experiment. If we maintain the values of temperature and density,

but alter  $t$  and  $R$  by a factor of  $10^{-4}$ , all  $\xi$ 's will be reduced by this factor. The  $\xi$ 's will then be at most of order 1. They drop below unity at a density  $\sim 10^{18} \text{ cm}^{-3}$ . Thus, the charge state freezes at  $10 > z > 6$ , as the plasma falls out of equilibrium. The scaling of  $\xi$ , by itself, prevents recombination of the plasma down to  $z \sim 1$ , as in a HANE.

Now, we consider an idealized laser experiment, shown in Table 2 and defined as follows. Starting with the HANE parameters of Table 1, scale time and expansion radius by  $10^{-4}$ . Density drops as in HANE according to a spherical expansion with constant velocity. Assume, however, that temperature is maintained at 500 eV to about 5 nsec. Thereafter, it drops adiabatically. The constant high temperature is due to the continued laser energy deposition. This provides a rough approximation to a laser experiment.

TABLE 2

| T<br>(ev) | $n$<br>( $\text{cm}^{-3}$ ) | $t$<br>(sec) | R<br>(cm) | $\xi_{R3}$ | $\xi_{RR}$ | $\xi_{RD}$ | $\xi_I$ | z  |
|-----------|-----------------------------|--------------|-----------|------------|------------|------------|---------|----|
| 5.0E+02   | 1.0E+21                     | 4.6E-10      | 4.6E-02   | 6.1E-02    | 3.5E+00    | 7.4E-01    | 1.6E+02 | 11 |
| 5.0E+02   | 1.0E+20                     | 1.0E-09      | 1.0E-01   | 1.3E-03    | 7.5E-01    | 1.6E-01    | 3.4E+01 | 11 |
| 5.0E+02   | 1.0E+19                     | 2.2E-09      | 2.2E-01   | 2.8E-05    | 1.6E-01    | 3.4E-02    | 7.3E+00 | 11 |
| 5.0E+02   | 1.0E+18                     | 4.6E-09      | 4.6E-01   | 6.1E-07    | 3.5E-02    | 7.4E-03    | 1.6E+00 | 11 |
| 1.1E+02   | 1.0E+17                     | 1.0E-08      | 1.0E+00   | 3.4E-06    | 2.7E-02    | 8.6E-07    | 7.6E-03 | 11 |
| 2.3E+01   | 1.0E+16                     | 2.2E-08      | 2.2E+00   | 2.8E-04    | 2.1E-02    | 0.0E+00    | 2.8E-10 | 11 |
| 5.0E+00   | 1.0E+15                     | 4.6E-08      | 4.6E+00   | 1.8E-02    | 1.6E-02    | 0.0E+00    | 0.0E+00 | 11 |
| 1.1E+00   | 1.0E+14                     | 1.0E-07      | 1.0E+01   | 2.0E-01    | 1.2E-02    | 0.0E+00    | 0.0E+00 | 11 |
| 2.3E-01   | 1.0E+13                     | 2.2E-07      | 2.2E+01   | 1.2E+00    | 9.6E-03    | 0.0E+00    | 0.0E+00 | 11 |

The first table entry shown is for an ion density of  $10^{21} \text{ cm}^{-3}$ . Maximum heating takes place at electron densities  $\sim 10^{21} \text{ cm}^{-3}$ . Although the laser does not penetrate to higher electron densities thermal conduction maintains the high temperature in the first table entry.

From Table 2 the plasma is in equilibrium at a temperature of 500 eV with  $z \geq 11$  as the dominant charge state. When the temperature begins to drop, at densities  $\leq 10^{18} \text{ cm}^{-3}$ , the  $\xi$ 's for recombination are less than unity. The expansion has reduced densities too much to allow for effective recombination at these temperatures. Eventually, when the temperature drops to very low values ( $< 1 \text{ eV}$ ) corresponding to a density  $\sim 10^{13} \text{ cm}^{-3}$ , 3-body recombination would reduce the  $z$  value of the debris. Once again, in the real laser experiment recombination will release energy to the electrons, maintaining a higher temperature and suppressing recombination. Furthermore, in the real laser experiment an ambient background gas will stop the expansion, halting the temperature decline. From that point the operative scaling becomes that of Eq. 1. We note, finally, that if in Table 2 we maintained the temperature and density values but scaled  $t$  and  $R$  by a factor of  $10^4$ , the  $\xi$ 's would be increased by that factor. Clearly, recombination to lower charge states would rapidly ensue. A HANE size laser experiment would recombine.

#### IV. Discussion and Conclusion

We compare the above results to a HANEX code simulation of the laser experiment. The state of aluminum target ions without the effects of background coupling was obtained by running the HANEX code with a background nitrogen density of  $10^{-6} \text{ Torr}$ . The target contained 40 cells, 13 of which blow off the front side. The initial conditions were chosen to match the series of shots described in the experimental coupling study of

Ripin, et al.<sup>(6)</sup> The target was 5.6 micron thick aluminum foil. The laser pulse length was 10 nsec with a full width half maximum of 4 nsec. The nominal laser energy was 100 joules with 50 joules within a radius of 125 microns. This produced a forward debris mass of 0.2 micrograms in a cone of half angle 40 degrees. In Table 3 we present the time history of a representative cell with an outward velocity of  $5.2 \times 10^7$  cm/sec. For comparison  $\xi_R = \xi_{RR} + \xi_{RD} + \xi_{R3}$  and  $\xi_I$  were calculated using the same rates as used for Table 2. The average charge state,  $\bar{z} = N_e/n$ , is given.

TABLE 3

| t<br>(nsec) | R<br>(cm) | n<br>(cm <sup>-3</sup> ) | N <sub>e</sub><br>(cm <sup>-3</sup> ) | T<br>(ev) | T <sub>e</sub><br>(ev) | $\xi_R$ | $\xi_I$ | $\bar{z}$ |
|-------------|-----------|--------------------------|---------------------------------------|-----------|------------------------|---------|---------|-----------|
| 3.38        | -0.002    | 6.5E+20                  | 7.0E+21                               | 200.      | 211.                   | 1.0E+02 | 1.6E+02 | 10.8      |
| 3.90        | 0.002     | 1.5E+20                  | 1.7E+21                               | 453.      | 489.                   | 5.5E+00 | 1.9E+02 | 11.1      |
| 4.42        | 0.012     | 5.8E+19                  | 6.5E+20                               | 482.      | 573.                   | 2.2E+00 | 1.0E+02 | 11.2      |
| 4.91        | 0.026     | 1.9E+19                  | 2.1E+20                               | 382.      | 590.                   | 7.9E-01 | 3.8E+0  | 11.3      |
| 5.38        | 0.044     | 7.1E+18                  | 8.0E+19                               | 293.      | 570.                   | 3.3E-01 | 1.5E+01 | 11.3      |
| 5.85        | 0.062     | 3.5E+18                  | 3.9E+19                               | 235.      | 550.                   | 1.8E-01 | 7.8E+00 | 11.3      |
| 6.55        | 0.097     | 1.4E+18                  | 1.5E+19                               | 170.      | 478.                   | 8.6E-02 | 2.9E+00 | 11.3      |
| 8.46        | 0.174     | 3.8E+17                  | 4.3E+18                               | 105.      | 269.                   | 4.1E-02 | 4.0E-01 | 11.3      |
| 11.9        | 0.384     | 5.9E+16                  | 6.6E+17                               | 48.       | 76.                    | 2.6E-02 | 8.7E-04 | 11.2      |
| 31.0        | 1.441     | 1.6E+15                  | 1.7E+16                               | 6.        | 7.                     | 3.7E-02 | 1.1E-24 | 11.2      |
| 54.0        | 2.699     | 2.5E+14                  | 2.8E+15                               | 2.        | 2.                     | 2.2E-01 | 1.7E-25 | 11.2      |

At early times when the laser pulse is still on, the ion and electron densities fall due to expansion but the electron temperature stays high first because the laser energy is being absorbed, and then, because of electron thermal conduction from the region of the target that is still

being heated. These effects were accounted for in Table 2 by assuming a constant temperature of 500 ev during the laser pulse. The debris expansion velocity is about a factor of 2 smaller than that of Table 2. This only changes the time scale for expansion by that amount, so Table 3 can be compared directly if time (t) and the  $\xi$ 's in Table 2 are increased by a factor of 2. We can see that after the laser is turned off (~ 10 nsec) the forward moving debris expands and cools adiabatically, as in our models.

The HANEX code includes laser absorption, radiation transport, time dependent chemistry, thermal conduction, etc. The agreement between code results and the simple models presented in Table 2 after the end of the laser pulse demonstrates that the controlling factors after 10 nsec are, in fact, adiabatic expansion and the recombination rates we discussed.

In conclusion, the simple model calculations given in Section III illustrate the essential differences between HANE and the laser experiment: a low debris charge state in the former and a high charge state in the latter. The primary effect follows from the breakdown of density scaling in the early debris expansion. A secondary effect is a result of the long deposition time of the laser energy, maintaining a higher temperature in the laser experiment than in a HANE at the same densities, even though at earliest times HANE temperatures are higher. We note that the results are essentially unchanged if we reduce this temperature even by a factor of 2 or 3. Recall, that even a scaled down HANE, in which the temperatures are an order of magnitude lower at corresponding densities still resulted in  $10 > z > 6$ . That is, the primary effect is the debris density scaling breakdown. Finally, we may ask to what extent charge exchange with an ambient background will reduce the debris charge state. We will present detailed simulation results in a separate report. However, we believe that

at high background density (0.5 - 5. Torr), where mixing of debris and background is limited, high charge states will persist. At lower background density (0.01 - 0.5 Torr) charge exchange may substantially reduce the charge states.

#### Acknowledgments

This work was supported by the Defense Nuclear Agency.

#### References

1. Clark, R. and V. Jacobs, unpublished.
2. B.H. Ripin, A.W. Ali, H.R. Griem, J. Grun, S.T. Kacenjar, C.K. Manka, E.A. McLean, A.N. Mostovych, S.P. Obenschain, J.A. Stamper, in Laser Interaction & Related Plasma Phenomena, Vol. 7, p. 857-877, ed. by H. Hora & G. Miley, Plenum 1986.
3. Giuliani, J.L. and M. Mul Brandon, "Numerical Simulations of the Laser-Target Interaction and Blast Wave Formation in the DNA/NRL Laser Experiment," NRL Memo No. 5762, 1986.
4. Longmire, C., M. Alme, R. Kilb, L. Wright, "Scaling of Debris-Air Coupling," MRC Report AMRC-R-338(1981).
5. See e.g., Zel'dovich and Raizer, "Physics of Shock Waves and High-Temperature Hydrodynamic Phenomena," Vol. I, Academic Press (1966), equation 6.104.
6. Ripin, B.H., E.A. McLean, J.H. Stamper, J. Grun, C.K. Manka, A.N. Mostovych, "Loss of Collisional Blast Wave Coupling with Pressure," ETHANL 7, 24 (1987).



## Appendix I

The rate coefficients for aluminum used to calculate the  $\xi$ 's for Tables 1 and 2 are given by the following expressions:

Recombination  $z \rightarrow z - 1$ :

$$\alpha_{RR} = A_R T_e^{-\eta}$$

$$\alpha_{RD} = A_D e^{-\left(\frac{E1_D}{T_e}\right)} \left[ 1 + B_D e^{-\left(\frac{E2_D}{T_e}\right)} \right] / T_e^{3/2}$$

$$\alpha_{R3} = \text{MAX}(\alpha_{L_{R3}}, \alpha_{H_{R3}}), \text{ where}$$

$$\alpha_{L_{R3}} = \frac{8.75 \times 10^{-27} z^3}{T_e^{9/2}} N_e \text{ MIN} \left[ 1, \frac{E_\infty}{100 T_e} \right] \quad (\text{low temperature})$$

$$\alpha_{H_{R3}} = \frac{N_e e^{\left(\frac{E_\infty}{T_e}\right)}}{6 \times 10^{21} T_e^{3/2}} \alpha_I \quad (\text{high temperature})$$

Ionization  $z - 1 \rightarrow z$ :

$$\alpha_I = \frac{A_I \times e^{-x}}{(x + B_I)} \quad , \quad \text{with } x = \frac{E_I}{T_e}$$

Here  $T_e$  is in ev, the  $\alpha$ 's are in  $\text{cm}^3/\text{sec}$  and  $E_\infty$  is the ionization energy of the  $z - 1$  ion in ev. The other constants are given in the following table.

| z  | $A_R$   | $\eta$  | $A_D$   | $B_D$   | $E1_D$  | $E2_D$  | $A_I$   | $B_I$   | $E_I$   |
|----|---------|---------|---------|---------|---------|---------|---------|---------|---------|
| 1  | 3.8E-13 | 7.9E-01 | 9.6E-10 | 2.0E-01 | 2.0E+00 | 6.0E-01 | 2.4E-06 | 0.0E+00 | 6.0E+00 |
| 2  | 2.1E-12 | 6.9E-01 | 2.5E-09 | 3.0E-01 | 2.9E+00 | 4.9E+00 | 2.8E-07 | 1.5E-01 | 1.9E+01 |
| 3  | 3.7E-12 | 8.2E-01 | 3.7E-09 | 0.0E+00 | 4.1E+01 | 0.0E+00 | 6.1E-08 | 2.3E-02 | 2.8E+01 |
| 4  | 8.7E-12 | 7.4E-01 | 6.2E-10 | 1.4E+01 | 2.8E+01 | 5.5E+01 | 8.0E-09 | 1.4E-01 | 1.2E+02 |
| 5  | 1.6E-11 | 7.2E-01 | 1.5E-09 | 1.1E+00 | 3.0E+01 | 7.7E+01 | 5.3E-09 | 1.3E-01 | 1.5E+02 |
| 6  | 2.4E-11 | 7.0E-01 | 2.6E-09 | 1.0E+00 | 3.1E+01 | 8.9E+01 | 2.6E-09 | 1.1E-01 | 1.9E+02 |
| 7  | 3.3E-11 | 6.9E-01 | 3.8E-08 | 8.0E-01 | 3.2E+01 | 1.2E+02 | 1.4E-09 | 1.2E-01 | 2.4E+02 |
| 8  | 4.9E-11 | 7.0E-01 | 2.2E-08 | 1.6E+01 | 2.7E+01 | 1.5E+02 | 7.0E-10 | 8.4E-02 | 2.8E+02 |
| 9  | 6.5E-11 | 7.1E-01 | 4.5E-08 | 5.2E+00 | 3.0E+01 | 1.7E+02 | 3.6E-10 | 3.0E-02 | 3.3E+02 |
| 10 | 1.1E-10 | 8.6E-01 | 1.3E-08 | 1.3E+01 | 1.8E+01 | 2.2E+02 | 2.3E-10 | 2.4E-01 | 4.0E+02 |
| 11 | 1.2E-10 | 8.3E-01 | 2.4E-08 | 0.0E+00 | 1.3E+03 | 0.0E+00 | 8.6E-11 | 2.1E-01 | 4.4E+02 |
| 12 | 1.8E-10 | 7.6E-01 | 7.9E-09 | 3.4E+01 | 2.5E+02 | 1.2E+03 | 5.3E-11 | 3.4E-01 | 2.1E+03 |
| 13 | 2.4E-10 | 7.5E-01 | 0.0E+00 | 0.0E+00 | 0.0E+00 | 0.0E+00 | 2.1E-11 | 3.4E-01 | 2.3E+03 |

In the HANEX code (but not in the calculations of  $\xi$  in Tables 1-3) the dielectronic recombination rates are reduced to account for density effects. Also, in the code only  $\alpha_{R3}^H$  is used for 3 body recombination because we do not encounter conditions in our high density background runs where  $\alpha_{R3}^L$  is important.

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